PRENATAL DIAGNOSIS

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The aim of the journal is to communicate the results of original research in a variety of clinical and scientific specialities concerned with *in utero* diagnosis of fetal abnormality in man (and animal models) resulting from genetic and environmental factors. This is considered to encompass genetic and other forms of screening aimed at identifying pregnancies at risk of fetal abnormality; the epidemiology and pathology of fetal abnormality (including fetal infection); fetal and maternal factors relating to intrauterine development; antenatal care in relation to the treatment and prevention of fetal abnormality; genetic counselling and selective termination of pregnancy; psychosocial aspects of prenatal diagnosis; the development and evaluation of services for prenatal diagnosis; developments in the field of obstetric ultrasound, fetoscopy and fetal blood sampling; the application of recombinant DNA technology and genetic linkage to the prenatal diagnosis of single gene defects; advances in amniotic cell culture techniques for the prenatal diagnosis of chromosome aberrations and metabolic defects.

EDITORIAL POLICY

The overriding criteria for publication are originality, a high scientific quality and interest to a wide audience of those concerned with all aspects of prenatal diagnosis research. The journal will provide a multidisciplinary forum for the exchange of information by which it is hoped to increase knowledge about the aetiology and pathogenesis of fetal abnormality and to promote further opportunities for its treatment and prevention. Papers not sufficiently substantiated by experimental detail will not be accepted, and although technical queries will be referred back to the author, the Editor reserves the right to make alterations in the text without altering the technical content.

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CONTENTS

VOLUME 4, ISSUE Nos. 1–6
and Special Issue

Issue No. 1, January–February

Prenatal Diagnosis of Chromosome Mosaicism *P. Benn, L. Y. F. Hsu, T. Perlis and A. Schonhaut* 1

Prenatal Diagnosis of Thalassaemia by Haemoglobin Chromatography on Biorex. An Evaluation of the Method *S. Dash, J. Panourgias, P. Karababa and D. Loukopoulos* 11

Normal Ultrasonic Size and Characteristics of the Fetal Adrenal Glands *P. Jeanty, F. Chervenak, P. Grannum and J. C. Hobbins* 21

Is Maternal Alpha-fetoprotein Screening Still of Value in a Low-risk Area for Neural Tube Defects? *J. G. Hooker, M. Lucas, B. A. Richards, I. M. Shirley, B. D. Thompson and R. H. T. Ward* 29

False Elevation of Amniotic Fluid Enzymes of Relevance for Prenatal Diagnosis: Creatine Kinase Measurement in Relation to Duchenne Muscular Dystrophy *R. J. Edwards, C. H. Rodeck and D. C. Watts* . 35

Activity and Multiple Forms of α-L-Fucosidase and Hexosaminidase in Chorion Biopsy Specimens and some Fetal Organs *E. M. Beyer and G. Ya. Wiederschain* 43

Discordance between Prenatal Cytogenetic Diagnosis and Outcome of Pregnancy *A. Loft and A. Tabor* 51

Prenatal Diagnosis of Fragile X in a Heterozygous Female Fetus and Postnatal Followup *M. G. Wilson and C. A. Marchese* 61

SHORT COMMUNICATIONS

Prenatal Diagnosis of Endocardial Fibroelastosis *L. Bovicelli, F. M. Picchio, G. Pilu, G. Baccarani, L. F. Orsini, N. Rizzo, G. Alampi, P. M. Benenati and J. C. Hobbins*. 67

Prenatal Diagnosis of Mosaicism 46,XX/46,XX,−21,+t, (21q21q) *W. S. Long, M. T. Mennuti, B. S. Emanuel and E. H. Zackai* . 73

SPECIAL REPORT

The Status of Fetoscopy and Fetal Tissue Sampling *M. S. Golbus* . 79

ANNOUNCEMENTS 83

Issue No. 2, March–April

Prenatal Diagnosis of Anhidrotic Ectodermal Dysplasia *M.-L. Arnold, R. Rauskolb, I. Anton-Lamprecht, A. Schinzel and W. Schmid* . . 85

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Mosaicism or Pseudomosaicism: The Problem of Hypermodal Cells in Amniotic Fluid Cell Culture *Y. Zhang, K. L. Garver, S. G. Marchese, G. R. Diggans, M. Muracca-Clemens and D. Flecker* 99

Evaluation of Inorganic Pyrophosphate in Amniotic Fluid as a Mode of Prenatal Diagnosis of Osteogenesis Imperfecta *K. L. Garver, M. G. Blitzer, G. Ibezim, S. G. Marchese, D. L. Pegram, A. M. Hagins and Y. Zhang* 109

Prenatal Diagnosis of Citrullinemia: Elevated Levels of Citrulline in the Amniotic Fluid in the Three Affected Pregnancies *W. J. Kleijer, W. Blom, J. G. M. Huijmans, M. C. T. Mooyman, R. Berger and M. F. Niermeijer* 113

Errors in Plasma Creatine Kinase Estimations on Fetal Blood Samples Resulting from Contamination with Amniotic Fluid and Maternal Blood: Relevance for the Prenatal Diagnosis of Duchenne Muscular Dystrophy *R. J. Edwards, C. H. Rodeck and D. C. Watts*. . . 119

SHORT COMMUNICATIONS

The Prenatal Diagnosis of Transient Cysts of the Fetal Choroid Plexus *P. Chudleigh, J. M. Pearce and S. Campbell* 135

Prenatal Diagnosis of Thrombocytopenia Absent Radius Syndrome using Ultrasound and Fetoscopy *K. Filkins, J. Russo, I. Bilinki, N. Diamond and B. Searle* 139

Prenatal Diagnosis of 45,X/46,XX Mosaicism: True Mosaicism in only One of Four Primary Cultures *L. A. Knight, H. Lawce, K. L. Wcislo and V. S. Bannister* 143

Amniotic Fluid Pregnancy-specific β1-glycoprotein (SP1) in Fetal Developmental Disorders *M. Heikinheimo, H. Jalanko, J. Leisti, K-L. Kolho, R. Salonen, H. Von Koskull and P. Aula* 147

Maternal Cell Contamination of Amniotic Fluid Cell Cultures from Two Consecutive Pregnancies Complicated by Fibroids *P. A. Benn, F. Gilbert and L. Y. F. Hsu* 151

Prenatal Diagnosis of Chondroectodermal Dysplasia with Fetoscopy *T-H. Bui, L. Marsk, O. Eklöf and K. Theorell* 155

LETTERS TO THE EDITOR 161

Issue No. 3, May–June

Transabdominal Fine Needle Biopsy from Chorionic Villi in the First Trimester *S. Smidt-Jensen and N. Hahnemann*. 163

The Effect of Oxygen Tension on Colony Formation and Cell Proliferation of Amniotic Fluid Cells *In Vitro* *K. R. Held and S. Sönnichsen* . 171

Experiences with Unexpected Structural Chromosome Aberrations in Prenatal Diagnosis in a Danish Series *L. O. Vejerslev and U. Friedrich* 181

Chemical Analysis of Succinylacetone and 4-hydroxyphenyllactate in Amniotic Fluid using Selective Ion Monitoring *C. Jakobs, L. Sweetman and W. L. Nyhan* 187

Prenatal Diagnosis of Hunter Syndrome *L. M. Archer, H. M. Kingston and P. S. Harper* 195

Prenatal Diagnosis of Mandibulofacial Dysostosis *K. H. Nicolaides, D. Johansson, D. Donnai and C. H. Rodeck* 201

Changes in Maternal Serum Alpha-fetoprotein Levels Associated with Intrauterine Genetic Diagnostic Procedures *G. F. Simpson, N-H. Sun and M. S. Golbus* 207

SHORT COMMUNICATIONS

Prenatal Diagnosis of Pericentric Inversion of Chromosome No 17 in a Twin Pregnancy *G. Kassam, A. T. L. Chen, M. F. Goldberg, S. Trusler and G. P. Oakley, Jr* 213

Mitomycin C Induced Chromosome Damage in Fetal Blood Cultures and Prenatal Diagnosis of Fanconi's Anaemia *J. Shipley, C. H. Rodeck, C. Garrett, J. Galbraith and F. Giannelli* 217

Genetic Amniocentesis in Multiple Gestations *K. Filkins, J. Russo, T. Brown, S. Schmerler and B. Searle* 223

Early Prenatal Ultrasonic Findings in Klippel–Trenaunay–Weber Syndrome *M. Seoud, R. Santos-Ramos and J. M. Friedman* . . . 227

A Rare Non-heterochromatic 9p+ Variant in Two Amniotic Fluid Cell Cultures *N. Archidiacono, V. Pecile, M. Rocchi, L. Dalprà, G. Nocera and G. Simoni* 231

Simple Ultrasonic Diagnosis of Osteogenesis Imperfecta Type II in Early Second Trimester *A. Ghosh, J. S. K. Woo, C. W. Wan and V. C. W. Wong* 235

Issue No. 4, July–August

Improved Direct Molecular Diagnosis and Rapid Fetal Sexing *D. I. Hoar, D. B. Haslam and D. M. Starozik* 241

Parental Response to Mid-trimester Therapeutic Abortion following Amniocentesis *O. W. Jones, N. E. Penn, S. Shuchter, C. A. Stafford, T. Richards, C. Kernahan, J. Gutierrez, P. Cherkin, S. Reinsch and B. Dixson* 249

Prenatal Detection of the Congenital Nephrotic Syndrome (Finnish Type) by Trehalase Assay in Amniotic Fluid *P. R. Morin, M. Potier, L. Dallaire and S. B. Melançon* 257

Amniotic Fluid Microvillar Enzyme Activities in the Early Detection of Fetal Abnormalities *D. J. H. Brock, D. Bedgood, C. Hayward, N. J. Carbarns and C. Gosden* 261

Creatine Kinase Estimation in Pure Fetal Blood Samples for the Prenatal Diagnosis of Duchenne Muscular Dystrophy *R. J. Edwards, D. C. Watts, R. L. Watts and C. H. Rodeck* 267

Chromosome Analysis of First Trimester Chorionic Villus Biopsies Prepared by a Maceration Technique *D. E. Heaton, B. H. Czepulkowski, D. H. Horwell and D. V. Coleman* 279

Selective Survival of Only the Healthy Fetus following Prenatal Diagnosis of Thalassaemia Major in Binovular Twin Gestation *A. Antsaklis, J. Politis, C. Karagiannopoulos, D. Kaskarelis, P. Karababa, J. Panourgias, M. Boussiou and D. Loukopoulos* 289

S-100 Protein in Amniotic Fluid of Anencephalic Fetuses *C. J. M. Sindic, M. Freund, N. Van Regemorter, Ch. Verellen-Dumoulin and P. L. Masson* 297

SHORT COMMUNICATIONS

Immunoreactive Trypsin Level in Fetoscopically-obtained Cord Sera of Second Trimester Fetuses *H. C. Ryley and C. H. Rodeck* . . 303

Prenatal Diagnosis of a 46,XX/47,XX,+12 Mosaic *P. K. A. Jensen, N. Hahnemann, H. Rehder, E. Schwinger and U. Friedrich* . . . 307

Amniotic Band Syndrome in Second Trimester associated with Fetal Malformations *K-G. Borlum* 311

LETTERS TO THE EDITOR 315

Issue No. 5, September–October

A Prospective Study of Amniotic Fluid Cholinesterases: Comparison of Quantitative and Qualitative Methods for the Detection of Open Neural Tube Defects *P. C. Wyvill, D. A. Hullin, G. H. Elder and K. M. Laurence* 319

Predictive Value of Amniotic Acetylcholinesterase Analysis in the Diagnosis of Fetal Abnormality in 3700 Pregnancies *D. A. Aitken, N. M. Morrison and M. A. Ferguson-Smith* 329

Maternal Serum Alpha Fetoprotein in Monozygotic and Dizygotic Twin Pregnancies *H. Thom, C. M. Buckland, A. G. M. Campbell, B. Thompson and V. Farr* 341

Prenatal Sex Diagnosis: Testosterone and FSH Levels in Mid-trimester Amniotic Fluids *D. Abeliovich, J. R. Leiberman, I. Teuerstein and J. Levy* 347

Gastroschisis and Omphalocele: Prenatal Diagnosis and Perinatal Management *J. M. Davidson, T. R. B. Johnson, Jr., D. T. Rigdon and B. H. Thompson* 355

Prenatal Diagnosis of Atypical Tay-Sachs Disease by Chorionic Villi Sampling *A. M. Besançon, J. P. Belon, L. Castelnau, Y. Dumez and L. Poenaru* 365

Cystic Hygromata in Trisomy 18 and 21 *J. M. Pearce, D. Griffin and S. Campbell* 371

Aneuploidy and Cystic Hygroma Detectable by Ultrasound *D. H. A. Redford, M. B. McNay, M. E. Ferguson-Smith and M. E. Jamieson* . 377

SHORT COMMUNICATIONS

Prenatal Diagnosis of Recurrence of Cystic Hygroma with Normal Chromosomes *B. Dallapiccola, L. Zelante, G. Perla and G. Villani* . 383

Prenatal Diagnosis of Trisomy 9 Mosaicism Possibly Limited to Fetal Membranes *R. A. Pfeiffer, R. Ulmer, A. Kniewald and E. Wagner-Thiessen* 387

The Prenatal Diagnosis of Joubert's Syndrome of Familial Agenesis of the Cerebellar Vermis *S. Campbell, C. Tsannatos and J. M. Pearce.* 391

Issue No. 6, November–December

Psychological Reactions to Fetoscopy: A Controlled Study *G. A. Fava, L. Michelacci, G. Trombini, L. Bovicelli and C. Orlandi* 397

Glial Fibrillary Acidic Protein in Amniotic Fluids from Pregnancies with Fetal Neural Tube Defects *M. Albrechtsen, E. Bock and B. Norgaard-Pedersen* 405

Trisomy 20 Mosaicism *P. Miny and I.-H. Pawlowitzki* 411

Prenatal Diagnosis of Cystic Fibrosis Using a Monoclonal Antibody Specific for Intestinal Alkaline Phosphatase *D. J. H. Brock, L. Barron, D. Bedgood and V. Van Heyningen* 421

Prenatal Assessment of Anterior Abdominal Wall Defects and Their Prognosis *L. Mann, M. A. Ferguson-Smith, M. Desai, A. A. M. Gibson and P. A. M. Raine* 427

SHORT COMMUNICATIONS

Early Prenatal Sonographic Appearance of Rare Thoraco-abdominal Eventration *J. W. Seeds, R. C. Cefalo, S. C. Lies and W. L. Koonyz* 437

Amniotic Band Disruption Syndrome Associated with Elevated Amniotic AFP and Normal Acetylcholinesterase Gel Test *D. A. Aitken, H. M. May, M. A. Ferguson-Smith, R. Howat and H. G. Kohler* . . 443

Amniotic Band Syndrome Causing Fetal Head Deformity *R. M. Hughes, R. J. Benzie and C. L. Thompson* 447

A Culture Vessel for Amniotic Fluid Cells Allowing Faster Preparation of Chromosome Slides *A. Tabor, A-M. Lind, A. Milbrat Andersen and J. Philip* 451

Prenatal Ultrasonic Findings in Congenital Chloride Diarrhoea *P. Kirkinen and P. Jouppila* 457

Pericentric Inversion of the Y Chromosome and Prenatal Diagnosis *L. R. Shapiro, R. O. Pettersen, P. L. Wilmot, D. Warburton, P. A. Benn and L. Y. F. Hsu* 463

Sonographic Diagnosis of Thoracic Gastroenteric Cyst *In Utero* *J. P. Newnham, J. V. Crues, III, A. L. Vinstein and A. L. Medearis* . . 467

LETTER TO THE EDITOR 473

Author Index 475

Key Word Index 477

Special Issue, Spring

Collaborative Studies in Prenatal
Diagnosis of Chromosome Aberrations

Edited by

J. L. Hamerton and M. A. Ferguson-Smith

Editorial *M. A. Ferguson-Smith* 1

Introduction *J. L. Hamerton* 3

Maternal Age Specific Rates for Chromosome Aberrations and Factors Influencing Them: Report of a Collaborative European Study on 52 965 Amniocenteses *M. A. Ferguson-Smith and J. R. W. Yates* . . 5

A Collaborative Study of the Segregation of Inherited Chromosome Structural Rearrangements in 1356 Prenatal Diagnoses *A. Boué and P. Gallano* 45

Outcome of Cases of *De Novo* Structural Rearrangements Diagnosed at Amniocentesis *D. Warburton* 69

Risk for Chromosome Abnormality at Amniocentesis following a Child with a Non-inherited Chromosome Aberration *J. Stene, E. Stene and M. Mikkelsen* 81

United States Survey on Chromosome Mosaicism and Pseudomosaicism in Prenatal Diagnosis *L. Y. F. Hsu and T. E. Perlis* 97

A Canadian Collaborative Study of Mosaicism in Amniotic Fluid Cell Cultures *R. G. Worton and R. Stern* 131

European Collaborative Study on Prenatal Diagnosis: Mosaicism, Pseudomosaicism and Single Abnormal Cells in Amniotic Fluid Cell Cultures *T.-H. Bui, L. Iselius and J. Lindsten* 145